

K884139 ACCI-GUARD(TM) REGULAR AND PEDIATRIC HOLDEROct 6, 1988
6 days to decisionK884139 · Product code: **FMI** · General Hospital
Source: <https://www.510kdatabase.net/k884139/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Needle, Hypodermic, Single Lumen (FMI)
Date received	Sep 30, 1988
Decision date	Oct 6, 1988
Days to decision	6 days
Third-party review	No

APPLICANT

Company	Medical Safety Products, Inc.
Location	Denver, CO, US
Contact	E SAGSTETTER
510(k) history	6 submissions · 6 cleared · 1988-1994

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k884139/>; Data sourced from FDA 510(k) public records (accessdata.fda.gov).510k Database is an independent platform not affiliated with the U.S. Food and Drug Administration. Always verify at accessdata.fda.gov. - Generated June 30, 2026